S. 2510

To amend the Public Health Service Act to provide revised standards for quality assurance in screening and evaluation of gynecologic cytology preparations, and for other purposes.

IN THE SENATE OF THE UNITED STATES

December 18, 2007

Ms. Landrieu (for herself and Mr. Isakson) introduced the following bill; which was read twice and referred to the Committee on Health, Education, Labor, and Pensions

A BILL

- To amend the Public Health Service Act to provide revised standards for quality assurance in screening and evaluation of gynecologic cytology preparations, and for other purposes.
 - 1 Be it enacted by the Senate and House of Representa-
 - 2 tives of the United States of America in Congress assembled,
 - 3 SECTION 1. SHORT TITLE.
 - 4 This Act may be cited as the "Cytology Proficiency
 - 5 Improvement Act of 2007".

1	SEC. 2. REVISED STANDARDS FOR QUALITY ASSURANCE IN
2	SCREENING AND EVALUATION OF
3	GYNECOLOGIC CYTOLOGY PREPARATIONS.
4	(a) In General.—Section 353(f)(4)(B)(iv) of the
5	Public Health Service Act (42 U.S.C. 263a(f)(4)(B)(iv))
6	is amended to read as follows:
7	"(iv) requirements that each clinical
8	laboratory—
9	"(I) ensure that all individuals
10	involved in screening and interpreting
11	cytological preparations at the labora-
12	tory participate annually in a con-
13	tinuing medical education program in
14	gynecologic cytology that—
15	"(aa) is approved by the Ac-
16	crediting Council for Continuing
17	Medical Education or the Amer-
18	ican Academy of Continuing
19	Medical Education; and
20	"(bb) provides each indi-
21	vidual participating in the pro-
22	gram with gynecologic cytological
23	preparations (in the form of ref-
24	erenced glass slides or equivalent
25	technologies) designed to improve

1	the locator, recognition, and in-
2	terpretive skills of the individual;
3	"(II) maintain a record of the cy-
4	tology continuing medical education
5	program results for each individual in-
6	volved in screening and interpreting
7	cytological preparations at the labora-
8	tory;
9	"(III) provide that the laboratory
10	director shall take into account such
11	results and other performance metrics
12	in reviewing the performance of indi-
13	viduals involved in screening and in-
14	terpreting cytological preparations at
15	the laboratory and, when necessary,
16	identify needs for remedial training or
17	a corrective action plan to improve
18	skills; and
19	"(IV) submit the continuing edu-
20	cation program results for each indi-
21	vidual and, if appropriate, plans for
22	corrective action or remedial training
23	in a timely manner to the laboratory's
24	accrediting organization for purposes
25	of review and on-going monitoring by

1	the accrediting organization, including
2	reviews of the continuing medical edu-
3	cation program results during on-site
4	inspections of the laboratory.".

- 5 (b) Effective Date and Implementation; Ter-6 mination of Current Program of Individual Pro-7 ficiency Testing.—
- 8 (1) Effective date and implementation.— 9 Except as provided in paragraph (2), the amend-10 ment made by subsection (a) applies to gynecologic 11 cytology services provided on or after the first day 12 of the calendar year beginning 1 year after the date 13 of the enactment of this Act, and the Secretary of 14 Health and Human Services (hereafter in this sub-15 section referred to as the "Secretary") shall issue 16 final regulations implementing such amendment not 17 later than 270 days after such date of enactment.
 - (2) TERMINATION OF CURRENT INDIVIDUAL TESTING PROGRAM.—The Secretary shall terminate the individual proficiency testing program established pursuant to section 353(f)(4)(B)(iv) of the Public Health Service Act (42 U.S.C. 263a(f)(4)(B)(iv)), as in effect on the day before the date of enactment of subsection (a), at the end of

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- 1 the calendar year which includes the date of enact-
- 2 ment of the amendment made by subsection (a).

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